

**Communicable Disease Epidemiology
and Immunization Section**

401 5th Avenue, Suite 1250
Seattle, WA 98104

206-296-4774 Fax 206-296-4803

TTY Relay: 711

www.kingcounty.gov/health

Health Advisory – CDC Recommends Increased Vigilance for Acute Flaccid Myelitis, 29 July 2016

Action requested:

- **Be aware that 21 confirmed cases of acute flaccid myelitis (AFM) have been reported to CDC during January 1–June 30, 2016 among persons 6 months to 64 years of age (median 7 years).**
- **Consider AFM in patients presenting with onset of acute focal limb weakness AND a magnetic resonance image (MRI) showing spinal cord lesion largely restricted to gray matter* AND spanning one or more spinal segments OR cerebrospinal fluid (CSF) showing pleocytosis (white blood cell count >5 cells/mm³)**
- **Collect specimens from patients suspected of having AFM as early as possible in the course of illness (preferably on the day of onset of limb weakness) including:**
 - CSF
 - Whole blood
 - Serum
 - Peripheral blood mononuclear cells (PBMC)
 - Stool
 - An NP aspirate, NP wash, or NP swab (with lower respiratory specimen if indicated)
 - An oropharyngeal swab

Collection of specimens as close to the onset of illness as possible has the best chance to yield a diagnosis.

- **Report confirmed or suspected cases of AFM promptly to Public Health at (206) 296-4774. Public Health will provide guidance on laboratory testing of specimens for enteroviruses, West Nile virus, and other infectious etiologies known to be associated with AFM.**
- **Please complete the patient summary form found here when reporting patients to Public Health: (<http://www.cdc.gov/acute-flaccid-myelitis/hcp/data.html>).**

Background: From January 1, 2016 through June 30, 2016, CDC received 36 reports of suspected acute flaccid myelitis (AFM) in persons from 20 U.S. states; a total of 21 met the Council of State and Territorial Epidemiologist (CSTE) case definition for a confirmed case of AFM and three were classified as probable. During the same period in 2015, CDC received only eight reports of suspected AFM, of which five were classified as confirmed. Dates of onset for confirmed cases ranged from December 1, 2015 through June 18, 2016; 48% (10/21) had onset of limb weakness after May 1, 2016. Because of the reports of a possible epidemiological association of EVD-68 and AFM in 2014, cerebrospinal fluid (CSF) specimens available from 86% (18/21) of confirmed cases were tested at CDC; all specimens were negative for enterovirus at CDC. Pleocytosis was present in 81% (17/21) of confirmed AFM cases with a median of 50/mm³ (range, 6-758/mm³).

Resources

- For more information on AFM: <http://www.cdc.gov/acute-flaccid-myelitis/index.html>
- For guidance on clinical management of patients with AFM: <http://www.cdc.gov/acute-flaccid-myelitis/downloads/acute-flaccid-myelitis.pdf>

Instructions for Completing the AFM Patient Summary Form

GENERAL. Clinicians should report all patients who meet the case definition (as specified on page 4) for AFM to their state or local health department using this *Patient Summary Form*.

- a. Clinicians should report patients who meet the case definition regardless of any laboratory results.
- b. This form should be completed by, or in conjunction with, a clinician who provided care to the patient during the neurologic illness.
- c. So that cases can be monitored in as real time as possible, this form should be submitted to the state or local health department as soon as possible after case identification.

CDC requests that state health departments also submit the *Patient Summary Form* to CDC to help monitor these cases at the national level. A form that is largely complete but has some information pending (e.g., hospital or health department laboratory results) or under investigation (e.g., polio vaccination history) should still be submitted as soon as possible, and the pending results can then be provided to CDC when they become available.

Demographics

1. **TODAY'S DATE.** Date that clinician is initiating completing the patient summary form.
2. **STATE ASSIGNED ID.** Alpha/numeric
3. **SEX.** Indicated whether the case-patient is male or female.
4. **DATE OF BIRTH.** Case-patient birth date.
5. **RESIDENCE.** State in which case-patient resides.
6. **COUNTY.** County in which case-patient resides.
7. **RACE.** Self-reported race of case-patient; more than one option may be reported.
8. **ETHNICITY.** Self-reported ethnicity of case-patient.
9. **DATE OF ONSET OF LIMB WEAKNESS.** Limb weakness onset date of case-patients.
10. **HOSPITALIZED?** Was case-patient hospitalized?
11. **DATE HOSPITALIZED.** Date case-patient FIRST hospitalized.
12. **DATE DISCHARGED.** Date case-patient discharged from LAST hospital (indicate if still hospitalized).
13. **DIED?** Did case-patient die from this illness?
14. **DATE OF DEATH.** Case-patient's date of death.

Signs/symptoms/condition at ANY time during the illness

If the answer to a question is truly UNKNOWN or no information is recorded in the medical record (NOT RECORDED or NR) then check the UNK/NR box, otherwise, leave answer blank.

15. **WHICH LIMBS HAVE BEEN ACUTELY WEAK?** Specify any/all limbs (arms and or legs) for which there was noted acute onset of focal weakness.
16. **DATE OF NEUROLOGIC EXAM.** The neurologic examination date recorded at most severe weakness to that point.

17. **REFLEXES IN THE AFFECTED LIMB(S).** Numeric value assigned to reflexes in affected limb(s) recorded at the most severe weakness to that point.
18. **SENSORY LOSS/NUMBNESS?** Has case-patient experienced any sensory loss or numbness in the affected limb(s) at any time during the illness?
19. **BURNING PAIN?** Has case-patient experienced any burning pain in the affected limb(s) at any time during the illness?
20. **SENSORY LEVEL ON THE TORSO?** Has case-patient experienced reduced sensation below a certain level below the torso at any time during the illness?
21. **CRANIAL NERVE FEATURES.** Did case-patient have any cranial nerve features? If YES, indicate the type experienced by the case-patient.
22. **BOWEL OR BLADDER INCONTINENCE?** Has case-patient experienced at any time during the illness bowel or bladder incontinence?
23. **CHANGE IN MENTAL STATUS?** Has case-patient experienced at any time during the illness a change in mental status?
24. **SEIZURES?** Has case-patient experienced any seizures at any time during the illness?
25. **RECEIPT OF POSITIVE PRESSURE VENTILATION?** Has case-patient received positive pressure ventilation, including invasive or non-invasive ventilation and BiPAP or CPAP?

Other patient information

26. **RESPIRATORY ILLNESS?** Did case-patient have a respiratory illness within the 4-week period before onset of limb weakness?
27. **RESPIRATORY ILLNESS ONSET DATE.** Case-patient's respiratory onset date.
28. **GASTROINTESTINAL ILLNESS?** Did case-patient have a gastrointestinal illness (e.g., diarrhea or vomiting) within the 4-week period before onset of limb weakness?
29. **GASTROINTESTINAL ILLNESS ONSET DATE.** Case-patient's gastrointestinal illness onset date.
30. **RASH?** Did case-patient have a new onset rash within the 4-week period before onset of limb weakness?
31. **RASH ONSET DATE.** Case-patient's rash onset date.
32. **FEVER?** Did case-patient have a fever ($\geq 38^{\circ}\text{C}/100.4^{\circ}\text{F}$), measured by parent or provider, within the 4-week period before onset of limb weakness?
33. **FEVER ONSET DATE.** Case-patient's fever onset date.
34. **IMMUNOSUPPRESSING AGENTS?** Did case-patient receive any immuno-suppressing agents within the 4-week period before onset of limb weakness?
35. **IF YES, LIST.** If any, list the date medication first administered, name of medication, how administered, and the dosage, duration, and overall amount received by case-patient.
36. **TRAVEL OUTSIDE U.S.?** Did case-patient travel outside the U.S. within the 4-week period before onset of limb weakness?
37. **IF YES, LIST.** If any, list the country(s) visited by the case-patient.
38. **UNDERLYING ILLNESSES?** Does the case-patient have any underlying illnesses?
39. **IF YES, LIST.** List the case-patient's underlying illness(es).
40. **FEVER ON DAY OF LIMB WEAKNESS ONSET?** Did the case-patient experience a fever (see definition in 32.) on the day of onset of limb(s) weakness?

Polio vaccination history

41. **IPV DOSES?** Indicate, if known, the number of documented inactivated polio vaccine doses received by the case-patient before the onset of limb weakness.
42. **OPV DOSES?** Indicate, if known, the number of documented oral polio vaccine doses received by the case-patient before the onset of limb weakness.
43. **DOCUMENTED POLIO VACCINE DOSES IF TYPE UNKNOWN?** If type of vaccine not known, indicate the total number of documented polio vaccine doses received by case-patient before the onset of weakness.

Neuroradiographic findings

44. **MRI OF SPINAL CORD PERFORMED?** Indicate whether case-patient had an MRI of the spinal cord performed.
45. **IF YES, NUMBER OF SPINAL MRIs PERFORMED.** If case-patient had spinal MRI performed, indicate the number of documented spinal MRIs performed.
For questions 46-71, complete based on results from the most abnormal MRI.
46. **DATE OF STUDY.** Date of the most abnormal MRI of the case-patient's spinal cord.
47. **LEVELS IMAGED.** Indicate the spinal cord levels imaged by MRI.
48. **LOCATION OF LESIONS.** Indicate the location of spinal cord lesions.
49. **CERVICAL CORD LEVEL.** Indicate whether the cervical level was affected.
50. **THORACIC CORD LEVEL.** Indicate whether the thoracic level was affected.
51. **AREAS OF SPINAL CORD AFFECTED.** For cervical and thoracic levels, indicate what spinal cord areas were affected.
52. **CORD EDEMA.** Was there cord edema?
53. **GADOLINIUM USED?** Was gadolinium used with the spinal cord MRI? *If NO, skip to question 59.*
54. **GRAY MATTER LESIONS.** For cervical/thoracic cord or conus lesions if gadolinium used, was there enhancement of any gray matter lesions?
55. **WHITE MATTER LESIONS.** For cervical/thoracic cord or conus lesions if gadolinium used, was there enhancement of any white matter lesions?
56. **CERVICAL/THORACIC NERVE ROOTS.** For cervical/thoracic cord or conus lesions if gadolinium used, was there enhancement of any cervical/thoracic nerve roots?
57. **VENTRAL NERVE ROOTS.** For cauda equina lesions if gadolinium used, was there enhancement of the ventral nerve roots?
58. **DORSAL NERVE ROOTS.** For cauda equina lesions if gadolinium used, was there enhancement of the dorsal nerve roots?
59. **BRAIN MRI PERFORMED.** Indicate whether case-patient had a brain/brainstem/cerebellum MRI performed.
If NO, skip to question 72.
60. **DATE OF STUDY.** Date of the MRI of the case-patient's brain.
61. **SUPRATENTORIAL LESIONS.** Were there any supratentorial lesions identified with the brain MRI?
62. **IF YES, INDICATE LOCATION.** Indicate location of supratentorial lesions identified with the brain MRI.
63. **BRAINSTEM LESIONS.** Were there any brainstem lesions identified with the brain MRI?
64. **IF YES, INDICATE LOCATION.** Indicate location of brainstem lesions identified with the brain MRI.

65. **CRANIAL NERVE LESIONS.** Were there any cranial nerve lesions identified with the brain MRI?
66. **IF YES, INDICATE CRANIAL NERVES.** Indicate in which cranial nerve(s) lesions were detected with the brain MRI.
67. **CEREBELLUM LESIONS.** Were there any lesions detected in the cerebellum?
68. **GADOLINIUM USED?** Was gadolinium used with the brain MRI? *If NO, skip to question 72.*
69. **SUPRATENTORIAL LESIONS.** If gadolinium used, was there enhancement of any supratentorial lesions?
70. **BRAINSTEM LESIONS.** If gadolinium used, was there enhancement of any brainstem lesions?
71. **CRANIAL NERVE LESIONS.** If gadolinium used, was there enhancement of any cranial nerve lesions?
72. **EMG DONE?** Indicate if an EMG was performed and if so, indicate the date.
73. **IF YES, ACUTE MOTOR NEUROPATHY?** If yes an EMG was done, was there evidence of acute motor neuropathy, motor neuropathy, motor nerve or anterior horn cell involvement?
74. **LUMBAR PUNCTURE PERFORMED?** Indicate if there was a CSF examination done (option for up to two. If more than 2 CSF examinations performed, list the first 2 performed.
 - 67a. **CSF from LP1.** Complete findings for lumbar puncture 1.
 - 67b. **CSF from LP2.** Complete findings for lumbar puncture 1.
75. **WAS CSF TESTED?** Complete information for CSF specimen testing for any of the pathogens listed, test type, test results, whether specimen was typed, and type if known. Provide information for the EARLIEST specimen collected if more than one CSF specimen collected and tested.
76. **WAS A RESPIRATORY TRACT SPECIMEN TESTED?** Complete information for respiratory tract specimen testing for any of the pathogens listed, test type, test results, whether specimen was typed, and type if known. Provide information for the EARLIEST specimen collected if more than one respiratory specimen collected and tested.
77. **WAS A STOOL SPECIMEN TESTED?** Complete information for stool specimen testing for any of the pathogens listed, test type, test results, whether specimen was typed, and type if known. Provide information for the EARLIEST specimen collected if more than one stool specimen collected and tested.
78. **WAS SERUM TESTED?** Complete information for serum specimen testing for any of the pathogens listed, test type, and test results. Provide information for the EARLIEST specimen collected if more than one serum specimen collected and tested.
79. **SPECIFIC ETIOLOGY?** Was/Is a specific etiology considered to be the most likely cause for the patient's neurological illness?
80. **IF YES, LIST.** List the etiology determined and reason(s) for considering it the most likely cause for the case-patient's neurological illness.
81. **SPECIMENS TO CDC.** If case-patient classified as confirmed or probable, will clinical specimens be sent to CDC for testing?
82. **SPECIMEN TYPES TO CDC.** If yes, indicate the specimen type(s) that will be sent to CDC.

Case Definition

In June 2015, the Council of State and Territorial Epidemiologists (CSTE) adopted a [standardized case definition for acute flaccid myelitis\[6 pages\]](#). As of August 1, 2015, a patient must meet the CSTE clinical criteria below to be considered either a confirmed or probable case of acute flaccid myelitis:

Acute Flaccid Myelitis case definition:

(<http://c.ymcdn.com/sites/www.cste.org/resource/resmgr/2015PS/2015PSFinal/15-ID-01.pdf>)

Clinical Criteria

An illness with onset of acute focal limb weakness AND

- a magnetic resonance image (MRI) showing spinal cord lesion largely restricted to gray matter and spanning one or more spinal segments, OR
- cerebrospinal fluid (CSF) with pleocytosis (white blood cell count >5 cells/mm³)

Case Classification***Confirmed:***

- An illness with onset of acute focal limb weakness AND
- MRI showing spinal cord lesion largely restricted to gray matter and spanning one or more spinal segments

Probable:

- An illness with onset of acute focal limb weakness AND
- CSF showing pleocytosis (white blood cell count >5 cells/mm³).

Acute Flaccid Myelitis: Patient Summary Form

FOR LOCAL USE ONLY

Name of person completing form: _____ State assigned patient ID: _____
 Affiliation _____ Phone: _____ Email: _____
 Name of physician who can provide additional clinical/lab information, if needed _____
 Affiliation _____ Phone: _____ Email: _____
 Name of main hospital that provided patient's care: _____ State: _____ County: _____

-----DETACH and transmit only lower portion to limbweakness@cdc.gov if sending to CDC-----

Acute Flaccid Myelitis: Patient Summary Form

Form Approved
OMB No. 0920-0009
Exp Date: 04/30/2016

Form to be completed by, or in conjunction with, a physician who provided care to the patient during the neurological illness. Once completed, submit to Health Department (HD). HD can also facilitate specimen testing.

1. Today's date ____/____/____ (mm/dd/yyyy)
2. State assigned patient ID: _____
3. Sex: ☐ M ☐ F
4. Date of birth ____/____/____
- Residence: 5. State _____ 6. County _____
7. Race: ☐ American Indian or Alaska Native ☐ Asian ☐ Black or African American ☐ Native Hawaiian or Other Pacific Islander ☐ White (check all that apply)
8. Ethnicity: ☐ Hispanic or Latino ☐ Not Hispanic or Latino
9. Date of onset of limb weakness ____/____/____ (mm/dd/yyyy)
10. Was patient admitted to a hospital? ☐ yes ☐ no ☐ unknown
11. Date of admission to **first** hospital ____/____/____
12. Date of discharge from **last** hospital ____/____/____ (or ☐ still hospitalized at time of form submission)
13. Did the patient die from this illness? ☐ yes ☐ no ☐ unknown
14. If yes, date of death ____/____/____

SIGNS/SYMPTOMS/CONDITION:				
	Right Arm	Left Arm	Right Leg	Left Leg
15. Since neurologic illness onset, which limbs have been acutely weak? [indicate yes(y), no (n), unknown (u) for each limb]	Y N U	Y N U	Y N U	Y N U
16. Date of neurologic exam (recorded at most severe weakness to point of completing this form) (mm/dd/yyyy)	____/____/____			
17. At the time of most severe weakness, reflexes in the most affected limb(s):	<input type="checkbox"/> Areflexic/hyporeflexic (0-1) <input type="checkbox"/> Normal (2) <input type="checkbox"/> Hyperreflexic (3-4+)			
At ANY time during the illness, was there:				
18. Any sensory loss/numbness in the affected limb(s), at any time during the illness? (paresthesias should not be considered here)	Y N U			
19. Any pain or burning in the affected limb(s)?	Y N U			
	Yes	No	Unk/Not Recorded (NR)	
20. Sensory level on the torso (i.e., reduced sensation below a certain level of the torso)?				
21. Did patient have any of the cranial nerve features below? (If yes, check all that apply):				
<input type="checkbox"/> Diplopia/double vision (If yes, circle the cranial nerve involved if known: 3 / 4 / 6)				
<input type="checkbox"/> Loss of sensation in face <input type="checkbox"/> Facial droop <input type="checkbox"/> Hearing loss <input type="checkbox"/> Dysphagia <input type="checkbox"/> Dysarthria				
22. Bowel or bladder incontinence?				
23. Change in mental status (e.g., confused, disoriented, encephalopathic)?				
24. Seizure(s)?				
25. Receipt of positive pressure ventilation, including invasive or non-invasive ventilation and including BiPAP or CPAP?				

Other patient information:

In the 4-weeks BEFORE onset of limb weakness, did patient:	Yes	No	Unk/NR	
26. Have a respiratory illness?				27. If yes, onset date ____/____/____
28. Have a gastrointestinal illness (e.g., diarrhea or vomiting)?				29. If yes, onset date ____/____/____
30. Have a new onset rash?				31. If yes, onset date ____/____/____
32. Have a fever, measured by parent or provider and $\geq 38.0^{\circ}\text{C}/100.4^{\circ}\text{F}$?				33. If yes, onset date ____/____/____

Public reporting burden of this collection of information is estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS D-74 Atlanta, Georgia 30333.

<p>34. Receive any immunosuppressing agent(s) (BEFORE WEAKNESS ONSET)?</p>				<p style="text-align: right;">Form Approved OMB No. 0920-0009 Exp Date: 04/30/2016</p> <p>35. If yes: Date of first administration: ____/____/_____ Name of medication: _____ Mode of administration: <input type="checkbox"/>IM <input type="checkbox"/>IV <input type="checkbox"/>Oral Dosage / duration / overall amount administered: _____</p>
<p>36. Travel outside the US?</p>				<p>37. If yes, list country:</p>
<p>38. At onset of limb weakness, does patient have any underlying illnesses?</p>				<p>39. If yes, list:</p>
<p>40. On the day of onset of limb weakness, did patient have a fever?</p>				<p>(see definition for fever above in 32.)</p>

Polio vaccination history:

<p>41. How many doses of inactivated polio vaccine (IPV) are documented to have been received by the patient before the onset of weakness?</p>	<p>_____ doses</p>	<p><input type="checkbox"/> unknown</p>
<p>42. How many doses of oral polio vaccine (OPV) are documented to have been received by the patient before the onset of weakness?</p>	<p>_____ doses</p>	<p><input type="checkbox"/> unknown</p>
<p>43. If you do not have documentation of the type of polio vaccine received what is total number of documented polio vaccine doses received before onset of weakness?</p>	<p>_____ doses</p>	<p><input type="checkbox"/> unknown</p>

Neuroradiographic findings:

MRI of spinal cord **44.** Was MRI of spinal cord performed? ☐ yes ☐ no ☐ unknown

45. If yes, how many documented spinal MRIs were performed? _____

*If yes to Q44, complete Q46-Q71 based on **most abnormal spine MRI*** **46.** Date of most abnormal spine MRI ____/____/____

47. Levels imaged: ☐cervical ☐thoracic ☐lumbosacral ☐unknown

<p>48. Location of lesions:</p>	<p><input type="checkbox"/>cervical cord <input type="checkbox"/>thoracic cord <input type="checkbox"/>conus <input type="checkbox"/>cauda equina <input type="checkbox"/>unknown</p>	<p>Levels of cord affected (if applicable):</p> <p>49. Cervical: _____ 50. Thoracic: _____</p>	
<p>For cervical and thoracic cord lesions</p>	<p>51. What areas of spinal cord were affected?</p>	<p><input type="checkbox"/>predominantly gray matter <input type="checkbox"/>predominantly white matter <input type="checkbox"/>both equally affected <input type="checkbox"/> unknown</p>	
	<p>52. Was there cord edema?</p>	<p><input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown</p>	
<p>53. Gadolinium (GAD) used: <input type="checkbox"/>yes <input type="checkbox"/>no <input type="checkbox"/> unknown <i>(If NO, skip to question 59)</i></p>			
<p>For cervical, thoracic cord or conus lesions</p>	<p>54. Did any gray matter lesions enhance with GAD?</p>	<p><input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown</p>	
	<p>55. Did any white matter lesions enhance with GAD?</p>	<p><input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown</p>	
	<p>56. Did any cervical / thoracic nerve roots enhance with GAD?</p>	<p><input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown</p>	
<p>For cauda equina lesions</p>	<p>57. Did the ventral nerve roots enhance with GAD?</p>	<p><input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown</p>	

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	58. Did the dorsal nerve roots enhance with GAD?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown
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MRI of brain

59. Was brain/brainstem/cerebellum MRI performed? ☐ yes ☐ no ☐ unknown (If NO, skip to Q72) **60.** Date of study ___/___/___

61. Any supratentorial (i.e. lobe, cortical, subcortical, basal ganglia, or thalamic) lesions	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
	62. If yes, indicate location(s)	<input type="checkbox"/> cortex <input type="checkbox"/> basal ganglia <input type="checkbox"/> thalamus <input type="checkbox"/> subcortex <input type="checkbox"/> unknown <input type="checkbox"/> Other (specify): _____
63. Any brainstem lesions?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
	64. If yes, indicate location:	<input type="checkbox"/> midbrain <input type="checkbox"/> pons <input type="checkbox"/> medulla <input type="checkbox"/> unknown
65. Any cranial nerve lesions?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
	66. If yes, indicate which CN(s):	CN___ <input type="checkbox"/> unilateral <input type="checkbox"/> bilateral CN___ <input type="checkbox"/> unilateral <input type="checkbox"/> bilateral CN___ <input type="checkbox"/> unilateral <input type="checkbox"/> bilateral CN___ <input type="checkbox"/> unilateral <input type="checkbox"/> bilateral
67. Any lesions affecting the cerebellum ?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
68. Gadolinium (GAD) used:	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	(If NO, skip to question 72)
69. Did any supratentorial lesions enhance with GAD?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
70. Did any brainstem lesions enhance with GAD?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	
71. Did any cranial nerve lesions enhance with GAD?	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	

72. Was an EMG done? ☐ yes ☐ no ☐ unknown If yes, date ___/___/___ (mm/dd/yyyy)

73. If yes, was there evidence of acute motor neuropathy, motor neuronopathy, motor nerve or anterior horn cell involvement? ☐ yes ☐ no ☐ unk

CSF examination: 74. Was a lumbar puncture performed? ☐ yes ☐ no ☐ unknown

If yes, complete 74 (a,b) (If more than 2 CSF examinations, list the first 2 performed)

	Date of lumbar puncture	WBC/mm3	% neutrophils	% lymphocytes	% monocytes	% eosinophils	RBC/mm3	Glucose mg/dl	Protein mg/dl
74a. CSF from LP1									
74b. CSF from LP2									

Pathogen testing performed:

75. Was CSF tested? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown		Specimen Collection Date ___/___/___	
If 'yes', was specimen tested for the following:			
<u>Enterovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	Test Type PCR	Test Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	Typed (if positive)? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> not done
<u>West Nile Virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	
<u>West Nile Virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	IgM	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate <input type="checkbox"/> Pending <input type="checkbox"/> Unknown	
<u>Herpes simplex virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	
<u>Cytomegalovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	
<u>Varicella zoster virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	
<u>Was other pathogen identified:</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	If positive for other pathogen, specify test type: _____	List other pathogen(s) identified: _____	

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76. Was a RESPIRATORY TRACT specimen tested? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown Specimen Collection Date __/__/____					
Type of specimen: <input type="checkbox"/> nasopharyngeal swab <input type="checkbox"/> nasal wash/aspirate <input type="checkbox"/> oropharyngeal swab <input type="checkbox"/> other, specify: _____					
If 'yes', was specimen tested for the following:					
<u>Enterovirus/rhinovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	Test Type PCR	Test Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	Typed (if positive)? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> not done	Type _____	
<u>Adenovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> not done	_____	
<u>Influenza virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	<input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> not done	_____	
<u>Was other pathogen identified:</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	If positive for other pathogen, specify test type: _____	List other pathogen(s) identified:			

77. Was a STOOL specimen tested? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown Specimen Collection Date __/__/____					
If 'yes', was specimen tested for the following:					
<u>Non-polio Enterovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	Test Type PCR	Test Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	Typed (if positive)? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> not done	Type _____	
<u>Poliovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	PCR	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending			
<u>Poliovirus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	Culture	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending			
<u>Was other pathogen identified:</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	If positive for other pathogen, specify test type: _____	List other pathogen(s) identified:			

78. Was SERUM tested? <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown Specimen Collection Date __/__/____					
If 'yes', was specimen tested for the following:					
<u>West Nile Virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	Test Type PCR	Test Result <input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Pending	Typed (if positive)?	Type	
<u>West Nile Virus</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	IgM	<input type="checkbox"/> Positive <input type="checkbox"/> Negative <input type="checkbox"/> Indeterminate <input type="checkbox"/> Pending <input type="checkbox"/> Unknown			
<u>Was other pathogen identified:</u> <input type="checkbox"/> yes <input type="checkbox"/> no <input type="checkbox"/> unknown	If positive for other pathogen, specify test type: _____	List other pathogen(s) identified:			

79. Was/Is a specific etiology considered to be the most likely cause for the patient's neurological illness? ☐ yes ☐ no ☐ unknown

80. If yes, please list etiology and reason(s) considered most likely cause _____

81. If patient is a confirmed or probable case, will specimens be sent to CDC for testing? ☐ yes ☐ no ☐ unknown

82. If yes, types of specimens that will be sent to CDC for testing:
☐ CSF ☐ Nasal wash/aspirate ☐ BAL spec ☐ Tracheal aspirate ☐ NP/OP swab ☐ Stool ☐ Serum ☐ Other, list _____

Acute Flaccid Myelitis case definition

(<http://c.ymcdn.com/sites/www.cste.org/resource/resmgr/2015PS/2015PSFinal/15-ID-01.pdf>)

Criteria

An illness with onset of acute focal limb weakness AND

- a magnetic resonance image (MRI) showing spinal cord lesion largely restricted to gray matter and spanning one or more spinal segments, OR
- cerebrospinal fluid (CSF) with pleocytosis (white blood cell count >5 cells/mm³)

Case Classification

Confirmed:

- An illness with onset of acute focal limb weakness AND
- MRI showing spinal cord lesion largely restricted to gray matter and spanning one or more spinal segments

Probable:

- An illness with onset of acute focal limb weakness AND
- CSF showing pleocytosis (white blood cell count >5 cells/mm³).